ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
AFCARS (Electronic Format)	52	2	2,972.89	309,077

Estimated Total Annual Burden Hours: 309,077.

Additional Information: Copies of the proposed collection maybe obtained 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: *grjohnson@acf,hhs.gov*.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the *Federal Register*. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Attn: Desk Officer for ACF, E-mail: *Katherine__T.__Astrick@omb.eop.gov*.

Dated: June 20, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–12515 Filed 6–23–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; National Institutes of Health Construction Grants—42 CFR Part 52b (Final Rule)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on December 7, 2004, pages 70697—70698, and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The NIH may not conduct or sponsor, and the respondent is not required to respond to, an information that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: National Institutes of Health Construction Grants—42 CFR Part 52b (Final Rule). *Type of Information Collection Request:* Extension of No. 0925–0424, expiration date 3/31/2005. *Need and Use of the Information Collection:* This request is for OMB review and approval of an extension for the information collection and recordkeeping requirements contained in the regulation codified at 42 CFR part 52b. The purpose of the regulation is to govern the awarding and administration of grants awarded by NIH and its components for construction of new buildings and the alteration, renovation, remodeling, improvement, expansion, and repair of existing buildings, including the provision of equipment necessary to make the buildings (or applicable part of the buildings) suitable for the purpose for which it was constructed. In terms of reporting requirements: Section 52b.9(b) of the regulation requires the transferor of a facility which is sold or transferred, or owner of a facility, the use of which has changed, to provide written notice of the sale, transfer or change within 30 days. Section 52b.10(f) requires a grantee to submit an approved copy of the construction schedule prior to the start of construction. Section 52b.10(g) requires a grantee to provide daily construction logs and monthly status reports upon request at the job site. Section 52b.11(b) requires applicants for a project involving the acquisition of existing facilities to provide the estimated cost of the project, cost of the acquisition of existing facilities, and cost of remodeling, renovating, or altering facilities to serve the purposes for which they are acquired. In terms of recordkeeping requirements: Section 52b.10(g) requires grantees to maintain daily construction logs and monthly status reports at the job site. *Frequency of Response:* On occasion. *Affected Public:* Non-profit organizations and Federal agencies. *Type of respondents:* Grantees. The estimated respondent burden is as follows:

ESTIMATED ANNUAL REPORTING AND RECORDKEEPING BURDEN

	Estimated annual number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total hour burden	Estimated total annual burden hours requested
Reporting:					
Section 52b.9(b)	1	1	.50	.50	.50
Section 52b.10(f)	(60)	1	1	60	60
Section 52b.10(g)	(60)	12	1	720	720
Section 52b.11(b)	100	1	1	100	100
Recordkeeping:					
Section 52b.10(g)	(60)	260	1	15,600	15,600
Total	101			16,481	16,481

The annualized cost to the public, based on an average of 60 active grants in the construction phase, is estimated at: $576,818. There are no Capital Costs to report. There are no operating or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information and recordkeeping are necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information and recordkeeping, including the methodology and assumptions used; (3) Enhance the quality, utility, and clarity of the information to be collected and the recordkeeping information to be maintained; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection and recordkeeping techniques of other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Office of Regulatory Affairs, New Executive Building, Room 10235, Washington, DC 20503, Attention Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Jerry Moore, NIH Regulations Officer, Office of Management Assessment, Division of Management Support, National Institutes of Health, 6011 Executive Boulevard, Room 601, MSC 7669, Rockville, Maryland 20852; call 301–496–4607 (this is not a toll-free number) or e-mail your request to *jm40z@nih.gov*.

Comments Due Date: Comments regarding this information collection and recordkeeping are best assured of having full effect if received on or before July 25, 2005.

Dated: June 17, 2005.

Jerry Moore,

Regulations Officer, National Institutes of Health.

[FR Doc. 05–12596 Filed 6–23–05; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Public Notice

AGENCY: Centers for Disease Control and Prevention (CDC), Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC), National Center for Infectious Disease (NCID), Division of Bacterial and Mycotic Diseases (DBMD) through its component Branches has lead technical responsibility for a number of Category A, B and C bioterrorism agents and their associated toxins (*Bacillus anthracis, Clostridium botulinum, Brucella sps., Burkholderia sps., Staphylococcus* entertoxin B, other food-or waterborne bacterial pathogens, and other bacterial agents). DBMD provides technical support for the Nation's prevention and control efforts for human anthrax disease. Since 2001, DBMD has been collecting anthrax immune plasma from Department of Defense volunteers who received the licensed Anthrax Vaccine Adsorbed (AVA) according to the licensed schedule. DBMD has contracted with industry to produce anthrax immune globulin (AIG) from the collected anthrax immune plasma using anion-exchange chromatography. Since 2003, DBMD has been evaluating the efficacy and pharmacokinetics of AIG in small animals. Preliminary results of these studies are now available, and are being released to the public domain to facilitate development of immunotherapeutic agents for treatment of human inhalational anthrax disease. DBMD will continue to conduct AIG studies in animals, and will release data to the public as soon as the results become available.

Persons or organizations who are interested in receiving the preliminary animal AIG study results, and in receiving future updates, should contact CDC and provide a mailing address.

CDC prefers to receive requests for data electronically. These requests can be e-mailed to the attention of Michael J. Detmer at *MDetmer@cdc.gov.* Mailed responses can be sent to the following address: Michael J. Detmer, Division of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, Centers for Disease Control and Prevention, 1600 Clifton Rd., NE., Mail Stop C–09, Atlanta, GA 30333.

FOR FURTHER INFORMATION CONTACT: *Technical:* Clare A. Dykewicz, M.D., M.P.H. Division of Bacterial and Mycotic Diseases, National Center for Infectious Diseases, Centers for Disease Control and Prevention (CDC), 1600 Clifton Rd. NE., Mail Stop C 09, Atlanta, GA 30333. Telephone (404) 639–4138, e-mail: *cad3@cdc.gov.*

Dated: June 13, 2005.

James D. Seligman,

Associate Director for Program Services, Centers for Disease Control and Prevention.

[FR Doc. 05–12497 Filed 6–23–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Alcohol Abuse and Alcoholism; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title U.S.C., as amended. The grant applications and the discussion could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute on Alcohol Abuse and Alcoholism Special Emphasis Panel ZAA1 HH (40) SPECIAL EMPHASIS PANEL REVIEW OF FELLOWSHIP APPLICATIONS.

Date: August 2, 2005.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Lorraine Gunzerath, PhD, MBA Scientific Review Administrator, National Institute on Alcohol Abuse and Alcoholism, Office of Extramural Activities, Extramural Project Review Branch, 5635 Fishers Lane, Room 3043, Bethesda, MD 20892–9304, 301–443–2369, *lgunzera@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants, National Institutes of Health, HHS)