proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions; (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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

For Further Information Contact: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Christy Thomsen, Director, Office of Communications and Public Liaison, NCCAM, 31 Center Drive, Room 2B11, Bethesda, MD 20892–2182; or fax your request to 301–402–4741; or e-mail *thomsenc@mail.nih.gov*. Ms. Thomsen can be contacted by telephone at 301–451–8876.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: February 14, 2006.

Christy Thomsen,

Director, Office of Communications and Public Liaison, National Center for Complementary and Alternative Medicine, National Institutes of Health.

[FR Doc. E6–2507 Filed 2–21–06; 8:45 am]

BILLING CODE 4167–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Statement of Organization, Functions, and Delegations of Authority

Part N, National Institutes of Health, of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services (HHS) (40 FR 22859, May 27, 1975, as amended most recently at 69 FR 64081, November 3, 2004, and redesignated from Part HN as Part N at 60 FR 56606, November 9, 1995), is amended as set forth below to reflect the reorganization of the National Human Genome Research Institute, Division of Intramural Research, by establishing (1) the Molecular Neurogenetics Section in the Medical Genetics Branch and (2) the Vascular Biology Section in the Genome Technology Branch. The sections are transferring from, respectively, the National Heart, Lung, and Blood Institute and the National Institute of Mental Health.

Section N–B, Organization and Functions, under the heading National Human Genome Research Institute (N4, formerly HN4), Division of Intramural Research (N45, formerly HN45) is amended as follows:

(1) In the Genome Technology Branch (N455, formerly HN455), immediately after the paragraph on Genomic Functional Analysis Section (N4556, formerly HN 4556), insert the following:

Vascular Biology Section (N4557, formerly HN 4557). Conducts clinical and laboratory investigations in the molecular mechanisms of cardiovascular disease including vascular cell biology, gene therapy, and cell cycle regulation of vascular cells.

(2) In the Medical Genetics Branch (N456, formerly HN456), immediately after the paragraph on Vertebrate Embryology Section (N4567, formerly HN4567), insert the following:

Molecular Neurogenetics Section (N4568, formerly HN 4568). (1) Conducts clinical and basic research into the factors contributing to the phenotypic variation observed in monogenic diseases, using Gaucher disease as a prototype disorder; (2) investigates the relationship between Gaucher disease and parkinsonism; and (3) explores new therapeutic approaches for Gaucher disease.

Delegations of Authority

All delegations and redelegations of authority to officers and employees of NIH that were in effect immediately prior to the effective date of this amendment and are consistent with this amendment shall continue in effect, pending further redelegation.

Dated: February 8, 2006.

Elias A. Zerhouni,

Director, National Institutes of Health.

[FR Doc. 06–1642 Filed 2–21–06; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Comments are invited on: (a) Whether the proposed collections of information are 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) ways to enhance 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.

Proposed Project: Mandatory Guidelines for Federal Workplace Drug Testing Programs (OMB NO. 0930–0158)—Revision

SAMHSA's Mandatory Guidelines for Federal Workplace Drug Testing Programs will request OMB approval for the Federal Drug Testing Custody and Control Form for Federal agency and federally regulated drug testing programs which must comply with the HHS Mandatory Guidelines for Federal Workplace Drug Testing Programs (69 FR 19644) dated April 13, 2004, and for the information provided by laboratories for the National Laboratory Certification Program (NLCP).

The Federal Drug Testing Custody and Control Form is used by all Federal agencies and employers regulated by the Department of Transportation to document the collection and chain of custody of urine specimens at the collection site, for laboratories to report results, and for Medical Review Officers to make a determination. The Federal Drug Testing Custody and Control Form approved by OMB three years ago is being resubmitted for OMB approval without any revision.

Prior to an inspection, a laboratory is required to submit specific information regarding its laboratory procedures. Collecting this information prior to an inspection allows the inspectors to thoroughly review and understand the laboratory's testing procedures before arriving at the laboratory.

The NLCP application form has not been revised compared to the previous form.

The annual total burden estimates for the Federal Drug Testing Custody and Control Form, the NLCP application, the NLCP inspection checklist, and NLCP recordkeeping requirements are shown in the following table.

Form/respondent	Burden/ response (hrs.)	Number of responses	Total annual burden (hrs.)
Custody and Control Form:			
Donor	.08	7,096,000	567,680
Collector	.07	7,096,000	496,720
Laboratory	.05	7,096,000	354,800
Medical Review Officer	.05	7,096,000	354,800
Laboratory Application	3.00	3	9
Laboratory Inspection Checklist	3.00	100	300
Laboratory Recordkeeping	250.00	50	12,500
Total			1,786,809

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 7–1044, One Choke Cherry Road, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: February 14, 2006.

Anna Marsh,

Director, Office of Program Services.

[FR Doc. 06–1597 Filed 2–21–06; 8:45 am]

BILLING CODE 4162–20–M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4912–N–17]

Notice of Availability of a Final Environmental Impact Statement for the Development of Stillwater Business Park, City of Redding, CA

AGENCY: Office of the Assistant Secretary for Community Planning and Development, HUD.

ACTION: Notice.

SUMMARY: HUD gives notice to the public, agencies, and Indian tribes that the City of Redding, CA, makes available to the public for comment the Final Environmental Impact Statement/ Final Environmental Impact Report (FEIS/FEIR) for the Stillwater Business Park project located in Redding CA. The City of Redding, CA has prepared the FEIS/FEIR under its authority as the Responsible Entity for compliance with the National Environmental Policy Act (NEPA) in accordance with 24 CFR 58.4, and under its authority as lead agency in accordance with the California Environmental Quality Act (CEQA). This notice is given in accordance with the Council on Environmental Quality regulations at 40 CFR parts 1500–1508. A HUD Economic Development Initiative (EDI) special purpose grant would be used for the project. Environmental Protection Agency (EPA) State and Tribal Assistance Grants (STAG) will also fund water and wastewater related infrastructure. EPA is acting as a cooperating agency for this process.

DATES: *Comments Due Date:* Comments due no later than March 24, 2006. Comments on the FEIS/FEIR should be addressed to the contact person listed below.

FOR FURTHER INFORMATION CONTACT: Nathan Cherpeski, City of Redding, 777 Cypress Ave., Redding, CA 96001, at (530) 225–4519 or *ncherpeski@ci.redding.ca.us.* The FEIS/ FEIR is available on the Internet and can be viewed or downloaded at: *http://ci.redding.ca.us/cm/major_pr/still_buspk.html.* Copies of the DEIS and Draft EIS/EIR are also available for viewing at the following locations:

City of Redding, Permit Center, 777 Cypress Ave., Redding, CA 96001.

City of Anderson Planning Department, 1887 Howard Street, Anderson, CA 96007.

Shasta County Library—Anderson Branch, 3200 West Center, Anderson, CA 96007.

Shasta County Department of Resource Management, Planning Division, 1855 Placer Street, Redding, CA 96001.

Shasta County Library, 1855 Shasta Street, Redding, CA 96001.

SUPPLEMENTARY INFORMATION: A Notice of Intent to prepare a draft EIS was published May 11, 2004. Scoping meetings were held on April 4, 2001, August 12, 2003, and June 2, 2004, to determine the issues for the EIS/EIR. A DEIS/DEIR was completed in May 2005. The DEIS/DEIR was the subject of public comments, both oral and written, provided by agencies, interested groups, and individuals, at a public hearing on April 12, 2005, and during the DEIS public comment period which extended from March 18, 2005, through May 2, 2005.

As a result of comments received and after meetings with EPA, USFWS, USACOE, and the California Department of Fish and Game, the City of Redding circulated a Supplemental Draft Environmental Impact Statement (SDEIS/DEIR) with a comment period from September 30, 2005, through November 14, 2005. A public open house was held October 26, 2005. Significant changes were made to the preferred alternative in the SDEIR/DEIR. Those changes are reflected in the preferred alternative described in the FEIS/FEIR. Developable acreage has been reduced and the size of the open space preserve has increased.

The preferred alternative is the development of a medium-to-large parcel business park through the acquisition of land, construction of major infrastructure components, and the provision of public services and utilities to serve the development. The City of Redding is proposing the development of the area east and northeast of the Municipal Airport in Redding, California. The proposed action study area is located on the *Enterprise and Cottonwood, California* 7.5-minute USGS quadrangles, Township 31 North, Range 4 West, Sections 2, 3, 10, 14, 15, 22, 23, 26, 34, and 35. A portion of the proposed location is classified as industrial and a portion as park under the Redding General Plan, adopted in 2000. The purpose and need for this project is to increase the activity of contributory economic sectors by constructing a medium to large parcel business park within the City of Redding sphere of influence capable of attracting and accommodating diverse business and industrial users.

The original proposal called for an approximate 687-acre business park consisting of 383 acres of developable land for a total of 4,410,400 sq. ft. of improvements for professional offices and industrial users. The preferred