Services, Lubbock, TX, from March 16 through March 25, 1994. The inspection also involved a concurrent investigation which included interviews with individuals knowledgeable about the firm's operations. The inspection and the investigation documented serious deviations from the applicable Federal regulations. Deviations identified in the inspection included, but were not limited to, the following: (1) Failure to collect blood by aseptic methods in a sterile system to protect against contamination (21 CFR 640.4(f)), in that: (a) current and former employees stated in affidavits that on numerous occasions employees broke the sterility barrier of blood containers and drained blood into vacutainer tubes or biohazard containers in order to conceal overbleeds, and (b) employees used an incorrect phlebotomy technique on numerous occasions thereby possibly contaminating the blood collection bags with room air; (2) failure to maintain records of donor adverse reaction reports (21 CFR 606.160(b)(1)(iii)), in that, on numerous occasions, employees did not document mild to moderate donor adverse reactions; (3) failure to follow standard operating procedures to adequately determine donor suitability (21 CFR 606.100(b)(1)), in that employees stated that: (a) donors were not always asked screening questions, such as high risk behavior questions, in order to expedite the donation process, (b) donors were sometimes asked if anything had changed since the last time they donated instead of being asked the required acquired immune deficiency syndrome (AIDS)-related behavior questions, and (c) individuals under the influence of alcohol were accepted as blood donors; and (4) failure to adequately and promptly notify the Director, Center for Biologics Evaluation and Research, of such errors or accidents in the manufacture of products that may affect the safety, purity or potency of any product pursuant to 21 CFR 600.14, in that, all known facts of the incidents involving a phlebotomist who used an incorrect phlebotomy technique, whereby the units may have become contaminated with room air, were not reported to the agency.

FDA determined that the deviations from Federal regulations were significant and constituted a danger to public health, warranting a suspension pursuant to 21 CFR 601.6(a). In a letter to Blood Systems, Inc., United Blood Services, dated June 6, 1994, FDA detailed the violations noted earlier in this document and suspended the firm's establishment and product licenses. In the same letter, FDA acknowledged receipt of letters dated April 13, April 15, and May 24, 1994, submitted by the firm in response to the Form FDA–483, Inspectional Observations, left at the close of the inspection. FDA concluded that the firm's promises of corrective action were not sufficient based on the seriousness of the documented deviations. It was FDA's view that the establishment and products failed to conform to applicable donor protection standards which are intended to ensure a continuous and healthy donor population, as well as standards designed to ensure the continued safety, purity, potency, and quality of products manufactured.

FDA's letter dated June 6, 1994, also stated that the agency's inspection and investigational findings, including evidence that records were knowingly falsified, demonstrated willfulness on the part of the firm. As a result, pursuant to 21 CFR 601.5(b) the firm was not given additional time to achieve compliance with the regulations. The same letter provided notice that FDA intended to initiate proceedings to revoke U.S. License No. 183–009 and product licenses issued to Blood Systems, Inc., United Blood Services pursuant to 21 CFR 601.5(b) and provided notice of opportunity for a hearing pursuant to 21 CFR 12.21(b). In a letter to FDA dated June 8, 1994, Blood Systems, Inc., voluntarily requested that its licenses for the Lubbock, TX, location be revoked and thereby waived its opportunity for a hearing. In a letter to the firm dated July 13, 1994, FDA acknowledged voluntary revocation of the establishment license (U.S. License No. 183–009) and the aforementioned product licenses of Blood Systems, Inc., United Blood Services at the Lubbock, TX, location only. In the July 13, 1994, letter to the firm, FDA restricted the interstate distribution of autologous units currently in inventory except in documented emergency situations, and permitted the firm to resume collections of allogeneic and autologous blood products intended for distribution within the State of Texas.

FDA has placed copies of letters relevant to the license revocations on file under the docket number found in brackets in the heading of this document in the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857. These documents are available for public examination in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

Accordingly, under section 351 of the Public Health Act (42 U.S.C. 262), 21 CFR 601.5, and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Director, Center for Biologics Evaluation and Research (21 CFR 5.68), the establishment license (U.S. License No. 183–009) and the product licenses issued to the Lubbock, TX, location of Blood Systems, Inc., United Blood Services for the manufacture of Whole Blood, Red Blood Cells, Plasma, Cryoprecipitated AHF, Platelets, and Source Leukocytes, were revoked, effective July 19, 1994.

This notice is issued and published under 21 CFR 601.8 and under authority delegated to the Director, Center for Biologics Evaluation and Research (21 CFR 5.67).

Dated: October 31, 1995.

Michael G. Beatrice,

Deputy Director, Center for Biologics Evaluation and Research.

[FR Doc. 95–29220 Filed 11–29–95; 8:45 am]

BILLING CODE 4160–01–F

Health Resources and Services Administration

Advisory Council on Nurses Education; Notice of Meeting Cancellation

In Federal Register Document 95–27189 appearing on page 55720 in the issue for Tuesday, November 2, 1995, the December 14–15, 1995, meeting of the ''National Advisory Council on Nurse Education and Practice'' will be cancelled.

Dated: November 27, 1995.

Jackie E. Baum,

Advisory Committee Management Officer, HRSA.

[FR Doc. 95–29260 Filed 11–29–95; 8:45 am]

BILLING CODE 4160–15–P

National Institutes of Health

National Institutes of Environmental Health Sciences; Amended Notice of Meeting

Notice is hereby given of the rescheduling of the meeting of the Environmental Health Sciences Review Committee, the notice of which was published in the Federal Register 60 FR 49848 on September 27, 1995. This meeting could not be convened on November 16–17 due to the partial shutdown of the Federal Government. it is rescheduled for December 3–5 at 6:00 p.m., at the Omni Europa Hotel, Chapel Hill, NC, and is closed to members of

the public on the same bases as provided in the initial notice.

Dated: November 28, 1995.

Susan K. Feldman,

Committee Management Officer, NIH.

[FR Doc. 95–29343 Filed 11–29–95; 8:45 am]

BILLING CODE 4140–01–M

Substance Abuse and Mental Health Services Administration (SAMHSA)

Cancellation of Receipt Date for SAMHSA Conference Grant Applications

AGENCY: Center for Substance Abuse Prevention and Center for Substance Abuse Treatment, SAMHSA.

ACTION: Cancellation of January 10, 1996 Application Receipt Date.

SUMMARY: Pending certainty on the fiscal year 1996 appropriation for SAMHSA, the Center for Substance Abuse Prevention (CSAP) and the Center for Substance Abuse Treatment (CSAT) are canceling the January 10, 1996, receipt date for applications for the following grant programs:

CSAP's Knowledge Dissemination Conference Grants (CFDA No. 93.174)

CSAT's Substance Abuse Treatment Conference Grants (CFDA No. 93.218)

For information regarding future receipt dates or for programmatic assistance, potential applicants should contact the following individuals:

CSAP: Ms. Luisa del Carmen Pollard, Division of Public Education and Dissemination, CSAP, Rockwall II Building, Suite 800, 5600 Fishers Lane, Rockville, Maryland 20857, Tele: (301) 443–0377.

CSAT: Ms. Nancy Kilpatrick, Office of Scientific Analysis and Evaluation, CSAT, Rockwall II Building, Suite 840, 5600 Fishers Lane, Rockville, Maryland 20857, Tele: (301) 443–8831.

Dated: November 24, 1995.

Richard Kopanda,

Acting Executive Officer, SAMHSA.

[FR Doc. 95–29261 Filed 11–30–95; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of the Assistant Secretary for Housing

[Docket No. FR–3917–N–30]

Notice of Proposed Information Collection for Public Comment

AGENCY: Office of the Assistant Secretary for Housing, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

DATES: Comments due date: January 29, 1996.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB Control Number and should be sent to: Oliver Walker, Housing, Department of Housing and Urban Development, 451—7th Street, SW., Room 9116, Washington, DC 20410.

FOR FURTHER INFORMATION CONTACT: Barbara D. Hunter, Telephone number (202) 708–3944 (this is not a toll-free number) for copies of the proposed forms and other available documents.

SUPPLEMENTARY INFORMATION: The Department will submit the proposed information collection to OMB for review, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35, as amended).

The Notice is soliciting comments from members of the public and affecting agencies concerning the proposed collection of information to: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions 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; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond; including through the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

This Notice also lists the following information:

Title of Proposal:

Section 8 Housing Assistance Payments Program, Additional Assistance Payments

Projects with HUD-held Mortgages 24 CFR Part 886, Subpart A, Housing Assistance Payment (HAP)

OMB Control Number: 2502–0407.

Description of the need for the information and proposed use: The Section 8 Housing Assistance Program, Part 886, Subpart A (Loan Management Set-Aside Special Allocations), authorized the use of Section 8 assistance in existing multifamily projects with HUD-insured or HUD-held mortgages, including Section 202 projects (except those receiving assistance under 24 CFR part 885) and projects sold by the Department subject to purchase money mortgages. The form HUD–52537, Section 8 HAP Contract (Part I), provides the administrative mechanism to obligate the necessary funds for the financially troubled projects aided under this regulation, a copy of which is attached.

It is also necessary to collect application information using the form HUD–52530 for the Department to evaluate applications for Section 8 Loan Management Set-Aside (LMSA) assistance.

Agency form numbers: HUD 52530 and 52537.

Members of affected public: Individuals, households and State/Local Government and Non-Profit Institutions.

An estimation of the total numbers of hours needed to prepare the information collection is 22,642, the number of respondents is 3,126, frequency of response is 1, and the hours of response is 7,243.

Status of the proposed information collection: Extension with change.

Authority: Section 3506 of the Paperwork Reduction Act of 1995, 44 U.S.C. Chapter 35, as amended.

Dated: November 7, 1995.

Nicolas P. Retsinas,

A/S Secretary for Housing—Federal Housing Commissioner.

[FR Doc. 95–29232 Filed 11–29–95; 8:45 am]

BILLING CODE 4210–27–M

Office of the Assistant Secretary for Housing-Federal Housing Commissioner

[Docket No. FR–3917–N–31]

Notice of Proposed Information Collection for Public Comment

AGENCY: Office of the Assistant Secretary for Housing-Federal Housing Commissioner, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

DATES: Comments due: January 29, 1996.