OGE's SES Performance Review Board as it was last published at 71 FR 71548–71549 (December 11, 2006).

Approved: September 12, 2007.

Robert I. Cusick,

Director, Office of Government Ethics.

The following officials have been selected as regular members of the SES Performance Review Board of the Office of Government Ethics:

Marilyn L. Glynn [Chair], General Counsel, Office of Government Ethics;

Daniel D. Dunning [Alternate Chair], Deputy Director for Administration and Information Management, Office of Government Ethics;

Rosalind A. Knapp, Deputy General Counsel, Department of Transportation;

Daniel L. Koffsky, Special Counsel, Office of Legal Counsel, Department of Justice; and

David Maggi, Chief, Ethics Law and Programs Division, Office of the Assistant General Counsel for Administration, Department of Commerce.

[FR Doc. E7–18518 Filed 9–19–07; 8:45 am]

BILLING CODE 6345–02–P

GOVERNMENT PRINTING OFFICE

Depository Library Council to the Public Printer; Meeting

The Depository Library Council to the Public Printer (DLC) will meet on Monday, October 15, 2007, through Wednesday, October 17, 2007, at Doubletree Hotel Crystal City, located at Arlington, Virginia. The sessions will take place from 8 a.m. to 5 p.m. Monday through Wednesday. The meeting will be held at the Doubletree Hotel Crystal City, 300 Army Navy Drive, Arlington, Virginia. The purpose of this meeting is to discuss the Federal Depository Library Program. All sessions are open to the public. The sleeping rooms available at the Doubletree Hotel Crystal City will be at the Government rate of $ 201.00 (plus applicable state and local taxes, currently 10.25%) a night for a single or double. The Doubletree Hotel Crystal City is in compliance with the requirements of Title III of the Americans With Disabilities Act and meets all Fire Safety Act regulations.

William H. Turri,

Acting, Public Printer of the United States.

[FR Doc. E7–18505 Filed 9–19–07; 8:45 am]

BILLING CODE 1520–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Re-allotment of FY 2006 Funds for the Low Income Home Energy Assistance Program (LIHEAP)

AGENCY: Office of Community Services, ACF, HHS.

ACTION: Notice of determination concerning funds available for reallotment.

C.F.D.A. Number: 93.568.

SUMMARY: In accordance with Section 2607(b)(1) of the Low Income Home Energy Assistance Act (the Act), Title XXVI of the Omnibus Budget Reconciliation Act of 1981 (42 U.S.C. 8621, *et seq.*), as amended, a notice was published in the **Federal Register** on August 1, 2007 announcing the Secretary's preliminary determination that $326,894 in Fiscal Year (FY) 2006 funds may be available for re-allotment. After a 30-day comment period, this amount has not changed. This notice announces that $326,894 will be re-allotted to current Low Income Home Energy Assistance Program (LIHEAP) grantees.

Pursuant to the statute cited above, funds will be re-allotted to LIHEAP grantees based upon the normal allocation formula as if the funds had been appropriated for FY 2007. No subgrantees or other entities may apply for these funds.

FOR FURTHER INFORMATION CONTACT: Nick St. Angelo, Director, Division of Energy Assistance, Office of Community Services, 370 L'Enfant Promenade, SW., Washington, DC 20447; telephone (202) 401–9351.

Dated: September 13, 2007.

Josephine B. Robinson,

Director, Office of Community Services.

[FR Doc. E7–18580 Filed 9–19–07; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0229]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices: Current Good Manufacturing Practice Quality System Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by October 22, 2007.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *baguilar@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0073. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance:

Medical Devices: Current Good Manufacturing Practice Quality System Regulations—21 CFR Part 820 (OMB Control Number 0910–0073)—Extension

Under section 520(f) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360j(f)), the Secretary of the Department of Health and Human Services (the Secretary) has the authority to prescribe regulations requiring that the methods used in, and the facilities and controls used for, the manufacture, pre-production design validation (including a process to assess the performance of a device but not including an evaluation of the safety and effectiveness of a device), packing, storage, and installation of a device conform to current good manufacturing practices (CGMPs), as described in such regulations, to assure that the device will be safe and effective and otherwise in compliance with the act.

The CGMP/Quality System (CGMP/QS) regulation implementing authority provided by this statutory provision is found under part 820 (21 CFR part 820) and sets forth basic CGMP requirements governing the design, manufacture, packing, labeling, storage, installation, and servicing of all finished medical