and from eligibility for, or involvement in, nonprocurement transactions (*e.g.*, grants and cooperative agreements) of the United States Government as defined in 45 CFR part 76 (Debarment Regulations) for a period of five (5) years, beginning on August 20, 2002;

(2) to exclude himself from serving in any advisory capacity to PHS, including but not limited to service on any PHS advisory committee, board, and/or peer review committee, or as a consultant for a period of five (5) years, beginning on August 20, 2002; and

(3) to submit a letter to the *Journal of Molecular and Cellular Cardiology* requesting retraction of Figure 1 in the article by Hui Liu, *et al., J. Mol. Cell. Cardiol.* 33:2001–2014, 2001, within 30 days of notification of this action. This requirement will be noted on the ALERT System until Dr. Yao sends a copy of the retraction letter to ORI.

FOR FURTHER INFORMATION CONTACT: Director, Division of Investigative Oversight, Office of Research Integrity, 5515 Security Lane, Suite 700, Rockville, MD 20852, (301) 443–5330.

Chris B. Pascal,

Director, Office of Research Integrity.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services

[Document Identifier: CMS–R–70]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare and Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA)), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* Information Collection Requirements in HSQ–110, Acquisition, Protection and Disclosure of Peer Review Organization Information and Supporting Regulations in 42 CFR, Sections 476.104, 476.105, 417.116, and 476.134.; *Form No.:* CMS–R–70 (OMB# 0938–0426); *Use:* The Peer Review Improvement Act of 1982 authorizes PROs to acquire information necessary to fulfill their duties and functions and places limits on disclosure of the information. These requirements are on the PRO to provide notices to the affected parties when disclosing information about them. These requirements serve to protect the rights of the affected parties.; *Frequency:* Reporting on occasion; *Affected Public:* Business or other for-profit, Individuals or households, Not-for-profit institutions.; *Number of Respondents:* 362; *Total Annual Responses:* 3729; *Total Annual Hours:* 60,919

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS's Web site address at *http://www.hcfa.gov/regs/prdact95.htm,* or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov*, or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the CMS Paperwork Clearance Officer designated at the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development and Issuances, Attention: Melissa Musotto, Room N2–14–26, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: August 28, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, Office of Strategic Operations and Strategic Affairs, Division of Regulations Development and Issuances.

[FR Doc. 02–22765 Filed 9–6–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0215]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Export of FDA Regulated Products—Export Certificates

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

DATES: Submit written comments on the collection of information by October 9, 2002.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Stuart Shapiro, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Mark L. Pincus, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1471.

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.

Export Certificates for FDA Regulated Products Under Sections 801(e) and 802 of the Federal Food, Drug, and Cosmetic Act—New Collection

FDA is requesting approval from OMB for the collection of information from the public associated with the export of FDA regulated products as indicated in sections 801(e) and 802 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 381(e) and 382), as amended.

In April 1996, a new law entitled ''The FDA Export Reform and Enhancement Act of 1996'' was enacted. It was designed to ease restrictions on exportation of unapproved products regulated by FDA and to facilitate such exportation by provide foreign governments certificates verifying that the products may be legally exported. Specifically, section 801(e)(4) of the act provides that persons exporting certain