submitted to the Office of Management and Budget (OMB) the following requirements for emergency review. We are requesting an emergency review because the collection of this information is needed before the expiration of the normal time limits under OMB's regulations at 5 CFR Part 1320(a)(2)(ii). This is necessary to ensure compliance with an initiative of the Administration. We cannot reasonably comply with the normal clearance procedures because of an unanticipated event, as stated in 5 CFR 1320.13(a)(2)(ii). The agency cannot reasonably comply with the normal clearance procedures because the application and user account registration form must have OMB clearance by September 2008 to meet the time necessary to begin CAS security administrator training and user account registration for new CROWNWeb alpha testers and CROWNWeb production users.

1. *Type of Information Collection Request:* New collection; *Title of Information Collection:* CROWNWeb Authentication Service (CAS) Account Form; *Form Number:* CMS–10210 (OMB#: 0938–NEW); *Use:* The CROWNWeb Authentication Service (CAS) application must be completed by any person needing access to the CROWNWeb system which include includes CMS employees, ESRD Network Organization staff and dialysis facilities staff. The CROWNWeb system is the system used as the collection point of data necessary for entitlement of ESRD patients to Medicare benefits and Federal Government monitoring and assessing of quality and type of care provided to renal patients. The data collected in CAS will provide the necessary security measures for creating and maintaining active CROWNWeb user accounts and collection of audit trail information required by the CMS Information Security Officers (ISSO). *Frequency:* Reporting—One-time; *Affected Public:* Business or other for-profit, Not-for-profit; *Number of Respondents:* 15,600; *Total Annual Responses:* 15,600; *Total Annual Hours:* 7,800.

CMS is requesting OMB review and approval of this collection by *August 29, 2008*, with a 180-day approval period. Written comments and recommendations will be considered from the public if received by the individuals designated below by *August 5, 2008.*

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS' Web Site address at *http://www.cms.hhs.gov/regulations/pra* or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov*, or call the Reports Clearance Office on (410) 786–1326.

In commenting on the proposed information collections please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in one of the following ways by *August 5, 2008:*

1. *Electronically.* You may submit your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, *Attention:* Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850
and,

OMB Human Resources and Housing Branch, *Attention:* CMS Desk Officer, New Executive Office Building, Room 10235, Washington, DC 20503, *Fax Number:* (202) 395–6974.

Dated: June 2, 2008.

Michelle Shortt,

Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. E8–12681 Filed 6–5–08; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0288]

Compliance Policy Guide Sec. 560.700 Processing of Imported Frozen Products of Multiple Sizes (e.g., Shrimp, Prawns, Etc.) (CPG 7119.10); Withdrawal of Guidance

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; withdrawal.

SUMMARY: The Food and Drug Administration (FDA) is announcing the withdrawal of Compliance Policy Guide Sec. 560.700 Processing of Imported Frozen Products of Multiple Sizes (e.g., Shrimp, Prawns, Etc.) (CPG 7119.10) (CPG Sec. 560.700). CPG Sec. 560.700 is included in FDA's Compliance Policy Guides Manual, which was listed in the Annual Comprehensive List of Guidance Documents that published on March 28, 2006.

DATES: The withdrawal is effective June 6, 2008.

FOR FURTHER INFORMATION CONTACT: Robert D. Samuels, Center for Food Safety and Applied Nutrition (HFS–325), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–2300.

SUPPLEMENTARY INFORMATION: In a notice containing a cumulative list of guidances available from the agency that published in the **Federal Register** on March 28, 2006 (71 FR 15422 at 15453), FDA included the Compliance Policy Guides Manual, which includes CPG Sec. 560.700. FDA is withdrawing CPG Sec. 560.700 because it is obsolete.

Dated: May 15, 2008.

Margaret O' K. Glavin,

Associate Commissioner for Regulatory Affairs.

[FR Doc. E8–12766 Filed 6–5–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial (PLCO) (NCI)

SUMMARY: In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial (PLCO). *Type of Information Collection Request:* REVISION (OMB #: 0925–0407, current expiry date 10/31/2008). *Need and Use of Information Collection:* This trial is designed to determine if screening for prostate, lung, colorectal and ovarian cancer can reduce mortality from these cancers which currently cause an estimated 254,900 deaths annually in the U.S. The design is a two-armed randomized trial of men and women aged 55 to 74 at entry. OMB first approved this study in 1993 and has approved it every 3 years since then through 2008. During the first approval period a pilot study was conducted to