of their qualifications (in particular, those that address the required qualifications, outlined above) and the current expertise needs of the Task Force. It is anticipated that 2 individuals will be invited to serve on the Task Force beginning in January, 2007. AHRQ will retain and consider for future vacancies the nominations of those not selected during this cycle.

ADDRESSES: Submit your response to: Helen Burstin, MD MPH, ATTN: USPSTF Nominations, Center for Primary Care, Prevention, and Clinical Partnerships, Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850.

Nomination Submissions

Nominations may submitted in writing or electronically, but must include (1) the applicant's current curriculum vitae, and (2) a letter explaining how this individual meets the qualification requirements and how he/she would contribute to the Task Force. The letter should also attest to the nominee's willingness to serve as a member of the Task Force.

AHRQ will later ask persons under serious consideration for membership to provide detailed information that will permit evaluation of possible significant conflicts of interest. Such information will concern matters such as financial holdings, consultancies, and research grants or contracts.

Nomination Selection

Nominations for the Task Force will be selected on the basis of qualifications as outlined above (see qualification requirements) and the current expertise needs of the Task Force.

Arrangement for Public Inspection

Nominations and applications are kept on file at the Center for Primary Care, Prevention and Clinical Partnerships, and are available for review during business hours. AHRQ does not reply to individual responses, but considers all nominations in selecting members. Information regarded as private and personal, such as a nominee's social security number, home and internet addresses, home telephone and fax numbers, or names of family members will not be disclosed to the public. This is in accord with agency confidentiality policies and Department regulations (45 CFR 5.67).

FOR FURTHER INFORMATION CONTACT:

Therese Miller at therese.miller@ahrq.hhs.gov.

SUPPLEMENTARY INFORMATION:

Background

Under Title IX of the Public Health Service Act, AHRQ is charged with enhancing the quality, appropriateness, and effectiveness of health care services and access to such services. AHRQ accomplishes these goals through scientific research and promotion of improvements in clinical practice, including prevention of diseases and other health conditions, and improvements in the organization, financing, and delivery of health care services (42 U.S.C. 299–299c–7 as amended).

The Task Force is an independent expert panel, first established in 1984 under the auspices of the U.S. Public Health Service, Currently, the USPSTF, under AHRQ's authorizing legislation (see in particular, 42 U.S.C. 299b-4(a)), is convened at the call of the Director of AHRQ. The Task Force is charged with rigorously evaluating the effectiveness, cost-effectiveness and appropriateness of clinical preventive services and formulating or updating recommendations for primary are clinicians regarding the appropriate provision of preventive services. The USPSTF transitioned to a standing Task Force in 2001. Current Task Force recommendations and associated evidence reviews are available on the Internet (http://

www.preventiveservices.ahrq.gov).

Dated: June 22, 2006.

Carolyn M. Clancy,

Director.

[FR Doc. 06–5782 Filed 6–28–06; 8:45 am] **BILLING CODE 4160–90–M**

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 71 FR 32349–32350, dated June 5, 2006) is amended to reflect the reorganization of the Office of the Chief Science Officer, Office of the Director, Centers for Disease Control and Prevention.

Section C–B, Organization and Functions, is hereby amended as follows:

Revise the functional statement for the *Office of the Chief Science Officer* (CAS), as follows:

After item (12), insert the following item: (13) provides oversight, training, monitoring, and quality assurance in the use of animals in research.

Delete item (10) of the functional statement for the *Scientific Resources Program (CVCE)*, *National Center for Infectious Diseases (CVC)*, and renumber the remaining items accordingly.

Dated: June 22, 2006.

William H. Gimson.

Chief Operating Officer, Centers for Disease Control and Prevention (CDC).

[FR Doc. 06–5785 Filed 6–28–06; 8:45 am]

BILLING CODE 4160-18-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N-0021]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Request for Samples and Protocols

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 July 31, 2006.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. 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.

FOR FURTHER INFORMATION CONTACT:

Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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.