

b. Current Budget Period Financial Progress.
 c. New Budget Period Program Proposed Activity Objectives.
 d. Budget.
 e. Measures of effectiveness.
 f. Additional requested information, including (1) data related to performance target goals; (2) data on progress toward achieving objectives; (3) an inventory of total individual capacity building assistance and proactive training for the reporting period; and (4) data related to the quality assurance system.

2. Second trimester interim progress report shall be due 30 days after the completion of the first eight (8) months of the project period. This second trimester progress report will serve as your non-competing continuation application for the next funding cycle. (See Continuing Application Requirements provided by Procurement and Grants Office.) This report must include elements a–f, as listed in the first trimester report, and be completed during this time period (months 5–8). The report should also include the following:

a. Base line and actual level of core performance indicators.
 b. Specific guidance, which will be provided by the CDC three months prior to the due date.

3. The third trimester progress report shall be due 30 days after the end of the budget period. This report must include elements a–f as listed in the first trimester report, elements a–b as listed in the second trimester report, and completed during this time period (months 9–12).

4. Financial status report is due no more than 90 days after the end of the budget period.

5. Final financial and performance reports are due no more than 90 days after the end of the project period.

These reports must be mailed to the Grants Management or Contract Specialist listed in the "Agency Contacts" section of this announcement.

VII. Agency Contacts

We encourage inquiries concerning this announcement.

For Pre-application Technical Consultation: Send questions regarding this application to DHAPCBAPT@CDC.GOV. You will receive a response within 24–48 hours.

For general questions, contact: Technical Information Management Section, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341. Telephone: 770-488-2700.

For program technical assistance, contact: Gerlinda Gallegos Somerville,

Public Health Analyst, Centers for Disease Control and Prevention, National Center for HIV, STD, and TB Prevention, Division of HIV/AIDS Prevention, Capacity Building Branch, 1600 Clifton Road, Mailstop E-40, Atlanta, GA 30333, Telephone: 404-639-2918. E-mail address: DHAPCBAPT@CDC.GOV.

For financial, grants management, or budget assistance, contact: Roslyn Curington, Grants Management Specialist, Centers for Disease Control and Prevention, Procurement and Grants Office, 2920 Brandywine Road, Room 3000, Atlanta, Georgia 30341-4146. Telephone: 770-488-2767, E-mail address: zlp8@cdc.gov.

VIII. Other Information

This and other CDC funding opportunity announcements can be found on the CDC Web site, Internet address: www.cdc.gov. Click on "Funding" then "Grants and Cooperative Agreements."

Dated: April 6, 2005.

William P. Nichols,

*Director, Procurement and Grants Office,
Centers for Disease Control and Prevention.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS-5033-N6]

Medicare Program; Cancellation of the April 13, 2005 Advisory Board Meeting on the Demonstration of a Bundled Case-Mix Adjusted Payment System for End-Stage Renal Disease Services

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Cancellation of meeting.

SUMMARY: This notice cancels the April 13, 2005 Advisory Board Meeting on the Demonstration of a Bundled Case-Mix Adjusted Payment System for End-Stage Renal Disease (ESRD) Services. We published the meeting notice in the **Federal Register** on March 25, 2005 (70 FR 15343).

DATES: *Effective Date:* The notice announcing the cancellation of the meeting is effective April 12, 2005.

FOR FURTHER INFORMATION CONTACT: Pamela Kelly by e-mail at ESRDAdvisoryBoard@cms.hhs.gov or telephone at (410) 786-2461.

SUPPLEMENTARY INFORMATION: On June 2, 2004, we published a **Federal Register**

notice requesting nominations for individuals to serve on the Advisory Board on the Demonstration of a Bundled Case-Mix Adjusted Payment System for End-Stage Renal Disease (ESRD) Services. The June 2, 2004 notice also announced the establishment of the Advisory Board and the signing by the Secretary on May 11, 2004 of the charter establishing the Advisory Board. On January 28, 2005, we published a **Federal Register** notice (70 FR 4132) announcing the appointment of eleven individuals to serve as members of the Advisory Board on the Demonstration of a Bundled Case-Mix Adjusted Payment System for ESRD Services, including one individual to serve as co-chairperson, and one additional co-chairperson, who is employed by CMS. The first public meeting of the Advisory Board was held on February 16, 2005. The second public meeting of the Advisory Board scheduled for April 13, 2005 has been cancelled.

Authority: 5 U.S.C. App. 2, section 10(a).

(Catalog of Federal Domestic Assistance Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: April 7, 2005.

Mark B. McClellan,

Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 05-7408 Filed 4-8-05; 1:51 pm]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Cooperative Agreement to Support the World Health Organization International Programme on Chemical Safety

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

I. Funding Opportunity Description

The Food and Drug Administration (FDA) is announcing its intent to accept and consider a single source application for the award of a cooperative agreement to the World Health Organization (WHO) to support the International Programme on Chemical Safety (IPCS). FDA anticipates providing \$90,000 (direct and indirect costs) in fiscal year 2005 in support of this project. Subject to the availability of Federal funds and successful performance, 2 additional years of support up to \$90,000 per year (direct and indirect costs) will be available. FDA will support the research