00007, Centers for Disease Control and Prevention (CDC), 2920 Brandywine Road, Room 3000, Atlanta, GA 30341, Telephone: (770) 488–2733, Email address: Vxm7@cdc.gov.

For program technical assistance, contact: Donald Ruberti, Senior Public Health Advisor, Prevention Services Research Branch, Division of HIV/AIDS Prevention, Centers for Disease Control and Prevention (CDC), 1600 Clifton Road, NE, Mail Stop E46, Atlanta, Georgia 30333, Telephone: (404) 639–2098, Email address: http://www.dor1@cdc.gov. See also the CDC Home Page on the Internet: http://www.cdc.gov

To receive additional written information and to request an application kit, call 1–888-GRANTS4 (1–888–472–6874). You will be asked to leave your name and address and will be instructed to identify the Announcement number of interest.

Dated: July 27, 1999.

John L. Williams,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–19675 Filed 7–30–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–0926]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Regulations Under the Federal Import Milk Act; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of July 26, 1999 (64 FR 40379). The document announced that a proposed collection of information has been submitted to the Office of Management and Budget for review and clearance under the Paperwork Reduction Act of 1995. The document was published with an inadvertent error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: LaJuana D. Caldwell, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 99–18927, appearing on page 40379 in the **Federal Register** of Monday, July 26, 1999, the following corrections are made:

1. On page 40379, in the first column, the Docket number is corrected to read ''99N–0926''; and in the second column, under the **SUPPLEMENTARY INFORMATION** caption, in the title of the proposed collection of information, the OMB control number ''0910–021'' is corrected to read ''0910–0212''.

Dated: July 27, 1999.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning and Legislation

[FR Doc. 99–19688 Filed 7–30–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

International Workshop on the Standardization of Whole Blood Coagulation Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing a workshop entitled ''International Workshop on the Standardization of Whole Blood Coagulation Devices.'' The focus of the workshop is to define the issues relating to the calibration of whole blood coagulation assays. Workshop participants will be asked to develop a proposal for standardizing the calibration of these devices. The proposal will be referred to a standards development organization.

DATE: The workshop will be held on August 13, 1999, 1 p.m. to 6 p.m.

ADDRESSES: The workshop will be held at the Washington Plaza Hotel, 10 Thomas Circle NW., Washington, DC 20005.

FOR FURTHER INFORMATION CONTACT: Sheila J. Murdock, Office of Surveillance and Biometrics (HFZ–510), Center for Devices and Radiological Health, Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–594–3060, FAX 301–594–2968, e-mail ''coagulation@cdrh.fda.gov''.

SUPPLEMENTARY INFORMATION: Whole blood clotting assays are used increasingly in the point of care testing environment. The calibration of these assays against plasma methods is achieved through a variety of approaches. Consequently, the consistency of results between different devices and the traceability of results to plasma methods are variable. Limited correlation between assays can be particularly problematic when monitoring anticoagulant drugs.

The workshop will focus on defining the issues relating to the calibration of whole blood coagulation devices. Workshop participants will collaborate on a proposal for the development of a standardized approach to the calibration of these assays. The proposal will be referred to a standards development organization.

In order to make the best use of limited workshop time, guest speakers will be asked to write a draft standardization proposal prior to the date of the workshop. This document will be posted on the CDRH website after July 15, 1999, at ''http://www.fda.gov/cdrh/meetings/coag.html''. Members of the public will be encouraged to e-mail comments and recommendations about this document to ''coagulation@cdrh.fda.gov''. Summaries of all e-mailed comments sent with author's name will be posted to the website in order to provide a forum for ongoing discussion up to the week of the workshop.

Those persons interested in attending the workshop should fax or e-mail their registration including name, title, affiliation (i.e., end-user, government nonregulatory, government regulatory, industry, professional organization, proficiency testing organization, trade press, standards development organization), mailing address, telephone number, fax number, e-mail address, and area of interest. There is no charge to attend the workshop, however, advance registration is requested due to limited seating. If you need special accommodations due to a disability, please contact Shirley L. Meeks at least 7 days in advance of the meeting, at the Office of Systems Management (HFZ–17), Center for Devices and Radiological Health, Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301–594–1283, ext. 105, FAX 301–827–2929, e-mail ''SLM@CRDH.FDA.GOV''.

Registration forms and the preliminary agenda may also be accessed at the CDRH website at ''http://www.fda.gov/cdrh/meetings/coag.html''. The workshop agenda includes presentations by guest speakers, small breakout group discussions and deliberation and refining of a standardization proposal. The final plenary session will include reports to the assembly from the smaller group discussions. Time will be provided for public comments at the end of this session. The draft standardization proposal will be finalized according to the

recommendations of workshop participants. A summary report of the workshop will be available on CDRH's website approximately 15 working days after the workshop. The CDRH home page may be accessed at ''http://www.fda.gov/cdrh''.

Dated: July 23, 1999.

Linda S. Kahan,

Deputy Director for Regulations Policy, Center for Devices and Radiological Health.

[FR Doc. 99–19689 Filed 7–30–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Drug Testing Advisory Board; Meeting

Pursuant to Public Law 92–463, notice is hereby given of the meeting of the Drug Testing Advisory Board of the Center for Substance Abuse Prevention in September 1999.

The first day (September 8) of the Drug Testing Advisory Board meeting will be closed from 8 a.m. until 4:30 p.m. and involves the review of sensitive National Laboratory Certification Program (NLCP) internal operating procedures and program development issues. Therefore, this portion of the meeting will be closed to the public as determined by the Administrator, SAMHSA, in accordance with 5 U.S.C. 552b(c)(2), (4), and (6) and 5 U.S.C. App.2, sec. 10(d).

The second day (September 9) of the Drug Testing Advisory Board meeting will be open from 8 a.m. until 3:30 p.m. The open session will include a roll call, general announcements, and a discussion of the information submitted by industry representatives regarding the use of alternative matrices (hair, sweat, oral fluids) and on-site tests to test for drugs of abuse. A public comment period will be scheduled during the open session. If anyone needs special accommodations for persons with disabilities please notify the Contact listed below.

An agenda for this meeting and a roster of board members may be obtained from: Ms. Giselle Hersh, Division of Workplace Programs, 5600 Fishers Lane, Rockwall II, Suite 815, Rockville, MD 20857, Telephone: (301) 443–6014.

Substantive program information may be obtained from the contact whose name and telephone number is listed below.

Committee Name: Drug Testing Advisory Board.

Meeting Date: September 8–9, 1999.

Place: Holiday Inn, 5520 Wisconsin Avenue, Chevy Chase, Maryland 20815.

Closed: September 8, 1999; 8 a.m.–4:30 p.m.

Open: September 9, 1999; 8 a.m. –3:30 p.m.

Contact: Donna M. Bush, Executive Secretary, Telephone: (301) 443–6014 and FAX: (301) 443–3031.

Sandi Stephens,

Committee Management Officer, Substance Abuse and Mental Health, Services Administration.

[FR Doc. 99–19687 Filed 7–30–99; 8:45 am]

BILLING CODE 4162–20–U

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Coastal Barrier Improvement Act of 1990; Amendments to the Coastal Barrier Resources System

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice.

SUMMARY: We, the Fish and Wildlife Service, have changed the boundaries on three maps of the Coastal Barrier Resources System, in Florida and South Carolina, as directed by Congress. In addition, recent legislation reactivates changes to eight maps in Florida that had been invalidated by a court decision on March 8, 1998. These changes are effective without the need to revise maps. We are using this notice to inform the public about the distribution and availability of the revised and reactivated maps.

DATES: The boundary revisions for these units became effective on October 21, 1998.

FOR FURTHER INFORMATION CONTACT: Dr. Benjamin N. Tuggle, Department of the Interior, U.S. Fish and Wildlife Service, Division of Habitat Conservation, (703) 358–2161.

SUPPLEMENTARY INFORMATION:

Background

In 1982, Congress passed the Coastal Barrier Resources Act (P.L. 97–348) to restrict Federal spending that could foster development of undeveloped coastal barriers along the Atlantic and Gulf of Mexico coasts. In the Coastal Barrier Improvement Act of 1990 (P.L. 101–591), Congress amended the CBRA to broaden the definition of a coastal barrier, and approved a series of maps entitled ''Coastal Barrier Resources System'' dated October 24, 1990. These maps identify and depict those coastal barriers located on the coasts of the Atlantic Ocean, Gulf of Mexico, and the Great Lakes, and in Puerto Rico and the Virgin Islands that are subject to the Federal funding limitations outlined in the CBRA.

The CBRA also defines our responsibilities regarding the System maps. We have official custody of these maps, and prepare and distribute copies of the maps. We published a notice of the filing, distribution, and availability of the maps entitled ''Coastal Barrier Resources System'' dated October 24, 1990, in the **Federal Register** on June 6, 1991, (56 FR 26304–26312). We have announced all subsequent map revisions in the **Federal Register**.

Section 101(e) of Public Law 105–277, enacted on October 21, 1998, requires us to revise the maps of Coastal Barrier Resources System Unit FL–35/FL–35P in Monroe County, Florida, and Unit SC–03 in Georgetown County, South Carolina. Section 134 of Public Law 105–277 requires us to revise the map of Unit M09 in Colleton County, South Carolina. Section 335 of Public Law 105–277 gives ''the force and effect of law'' to a set of eight maps entitled ''Coastal Barrier Resources System'', dated October 24, 1990, revised November 12, 1996, that were previously announced in the **Federal Register** on May 28, 1997 (62 FR 28891–28892). These eight maps were invalidated by a court order on March 5, 1998 (*Coastal Alliance* v. *Babbitt,* Civil Action No. 97–1344 (D. D.C.)), but were reinstated as of October 21, 1998, by Section 335 of Public Law 105–277.

Three Revised Maps in Florida and South Carolina

Consistent with Congressional instructions, the following three maps were modified:

North Key Largo Unit FL–35/FL–35P—Congress excluded Pumpkin Key from the System. Congress also removed some developed property of the Ocean Reef community from the System.

Huntington Beach Unit SC–03—Congress changed the western boundary of this unit to remove developed property from the System.

Edisto Island Unit M09—Congress changed the southern and western boundary of this unit back to the boundary it established in 1982.

Eight Reactivated Maps in Florida

We substituted the eight maps dated ''October 24, 1990, revised November 12, 1996,'' for the previous maps for the units. These maps are described in the May 28, 1997, **Federal Register** (62 FR 28891–28892) and pertain to the following units: