decision in part on a submission labeled CBI, then a nonconfidential version of the document that summarizes the key data or information should be submitted for the public docket. To ensure that proprietary information is not inadvertently placed in the docket, submissions containing such information should be sent directly to the contact person listed above and not to the public docket. Information covered by a claim of confidentiality will be disclosed by EPA only to the extent allowed and by the procedures set forth in 40 CFR part 2. If no claim of confidentiality accompanies the submission when EPA receives it, EPA will make it available to the public without further notice to the person making comments.

Dated: November 3, 2005.

Elizabeth Craig,

Acting Assistant Administrator, Office of Air and Radiation.

[FR Doc. 05–22995 Filed 11–18–05; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

Public Information Collection(s) Requirement Submitted to OMB for Emergency Review and Approval

November 16, 2005.

SUMMARY: The Federal Communications Commission, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s), as required by the Paperwork Reduction Act of 1995, Public Law 104–13. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number. Comments are requested concerning (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology.

DATES: Written Paperwork Reduction Act (PRA) comments should be submitted on or before December 21, 2005. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contacts listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Kristy L. LaLonde, Office of Management and Budget, Room 10234 NEOB, Washington, DC 20503, (202) 395–3087, or via fax at 202–395–5167 or via Internet at *Kristy_L._LaLonde@omb.eop.gov*, and Judith B. Herman, Federal Communications Commission, Room 1–C804, 445 12th Street, SW., Washington, DC 20554. You may submit your Paperwork Reduction Act (PRA) comments by e-mail or U.S. postal mail. To submit your comments by e-mail, send them to: *PRA@fcc.gov*. To submit your comments by U.S. mail, mark it to the attention of Judith B. Herman, Federal Communications Commission, 445 12th Street, SW., Room 1–C804, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection(s), send an e-mail to *PRA@fcc.gov* or contact Judith B. Herman at 202–418–0214.

SUPPLEMENTARY INFORMATION: The Commission has requested emergency OMB processing review of this new information collection with an OMB approval by November 18, 2005.

OMB Control Number: 3060–1004.

Title: Revision of the Commission's Rules to Ensure Compatibility with Enhanced 911 Emergency Calling Systems.

Form No.: N/A.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit, not-for-profit institutions, and state, local and tribal government.

Number of Respondents: 50 respondents; 213 responses.

Estimated Time Per Response: 5–200 hours.

Frequency of Response: Quarterly, semi-annual and one-time reporting requirements, third party disclosure requirement, and recordkeeping requirement.

Total Annual Burden: 1,202 hours.

Total Annual Cost: N/A.

Privacy Act Impact Assessment: N/A.

Needs and Uses: The Commission is seeking emergency OMB processing of this information collection by November 18, 2005. We have revised this information collection because on October 21, 2005, the Commission released an order (FCC 05–181) finding that certain Tier III carriers did not sufficiently support their requests for waiver of the E911 rules, but providing the carriers with additional time, until July 21, 2006, to augment the record to show a clear path to full compliance with the E911 requirements. The Commission also imposed conditions and required Tier III carriers to file separate status reports by November 21, 2005, and commencing February 1, 2006, additional status reports on a quarterly basis, for a two-year period.

In addition, on October 28, 2005 (FCC 05–182) and on November 3, 2005 (FCC 05–188), in response to requests for relief submitted by certain Tier III carriers, the Commission released orders that granted, in part, limited extensions of the December 31, 2005 requirement, subject to conditions, and required Tier III carriers to file status reports on a quarterly basis, for a two-year period beginning on February 1, 2006. Further, FCC 05–188 required one Tier III carrier, in addition to the quarterly reporting requirements, to submit a compliance plan by November 3, 2006.

The Commission will use the information submitted by Tier III carriers subject to reporting requirements to ensure that they comply with the Commission's E911 requirements and the terms of the underlying orders addressing their requests for waiver relief.

Federal Communications Commission.

Marlene H. Dortch,

Secretary.

[FR Doc. 05–23098 Filed 11–18–05; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Meeting of the President's Council on Bioethics on December 8–9, 2005

AGENCY: The President's Council on Bioethics, HHS.

ACTION: Notice.

SUMMARY: The President's Council on Bioethics (Edmund D. Pellegrino, MD, Chairman) will hold its twenty-second meeting, at which, among other things, it will discuss ethical issues relating to children. Subjects discussed at past Council meetings (though not on the agenda for the present one) include: Cloning, assisted reproduction, reproductive genetics, IVF, ICSI, PGD, sex selection, inheritable genetic modification, patentability of human organisms, neuroscience, aging

retardation, lifespan-extension, and organ procurement for transplantation. Publications issued by the Council to date include: *Human Cloning and Human Dignity: An Ethical Inquiry* (July 2002); *Beyond Therapy: Biotechnology and the Pursuit of Happiness* (October 2003); *Being Human: Readings from the President's Council on Bioethics* (December 2003); *Monitoring Stem Cell Research* (January 2004), *Reproduction and Responsibility: The Regulation of New Biotechnologies* (March 2004), *Alternative Sources of Human Pluripotent Stem Cells: A White Paper* (May 2005), and *Taking Care: Ethical Caregiving in Our Aging Society* (September 2005).

DATES: The meeting will take place Thursday, December 8, 2005, from 9 a.m. to 4:30 p.m. ET; and Friday, December 9, 2005, from 8:30 a.m. to 12:30 p.m. ET.

ADDRESSES: City Center Hotel, 1143 New Hampshire Avenue, NW., Washington, DC 20037. Phone 202–775–0800.

Agenda: The meeting agenda will be posted at *http://www.bioethics.gov*.

Public Comments: The Council encourages public input, either in person or in writing. At this meeting, interested members of the public may address the Council, beginning at 11:30 a.m., on Friday, December 9. Comments are limited to no more than five minutes per speaker or organization. As a courtesy, please inform Ms. Diane Gianelli, Director of Communications, in advance of your intention to make a public statement, and give your name and affiliation. To submit a written statement, mail or e-mail it to Ms. Gianelli at one of the addresses given below.

FOR FURTHER INFORMATION CONTACT: Ms. Diane Gianelli, Director of Communications, The President's Council on Bioethics, Suite 700, 1801 Pennsylvania Avenue, Washington, DC 20006. Telephone: 202/296–4669. E-mail: *info@bioethics.gov*. Web site: *http://www.bioethics.gov*.

Dated: November 16, 2005.

Richard Roblin,

Acting Executive Director, The President's Council on Bioethics.

[FR Doc. 05–22990 Filed 11–18–05; 8:45 am]

BILLING CODE 4154–06–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Fees for Sanitation Inspections of Cruise Ships

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: This notice announces an increase in the fees for vessel sanitation inspections for the remainder of fiscal year 2006 (January 1, 2006, through September 30, 2006).

DATES: *Effective* January 1, 2006.

FOR FURTHER INFORMATION CONTACT: David L. Forney, Chief, Vessel Sanitation Program, National Center for Environmental Health, Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, NE., Mailstop F–23, Atlanta, Georgia 30341–3724, telephone (770) 488–7333, E-mail: *Dforney@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose and Background

The fee schedule for sanitation inspections of passenger cruise ships inspected under the Vessel Sanitation Program (VSP) was first published in the **Federal Register** on November 24, 1987, (52 FR 45019), and CDC began collecting fees on March 1, 1988. Since 1987, CDC has published the fee schedule annually. The fees schedule for FY 2006 was initially announced in the **Federal Register** on September 7, 2005, with fees effective October 1, 2005. Because of a significant increase in travel expenses and personnel costs, VSP will be increasing inspection fees effective January 1, 2006. This will be the first increase in fees since FY 2002.

The formula used to determine the fees is as follows:

$$\text{Average cost per inspection} = \frac{\text{Total cost of VSP}}{\text{Weighted number of annual inspections}}$$

The average cost per inspection is multiplied by a size/cost factor to determine the fee for vessels in each size category. The size/cost factor was established in the proposed fee schedule published in the **Federal Register** on July 17, 1987, (52 FR 27060) and revised in a schedule published in the **Federal Register** on November 28, 1989, (54 FR 48942). The revised size/cost factor is presented in Appendix A.

Fee

The fee schedule (Appendix A) will be effective January 1, 2006, through September 30, 2006. If travel expenses continue to increase, the fees may need adjustment before September 30, 2006, because travel constitutes a sizable portion of VSP's costs. If an adjustment is necessary, a notice will be published in the **Federal Register** 30 days before the effective date.

Applicability

The fees will apply to all passenger cruise vessels for which inspections are conducted as part of CDC's VSP.

Dated: November 7, 2005.

Carlton Duncan,

Deputy Chief Operating Officer, Office of the Director, Centers for Disease Control and Prevention (CDC).

Appendix A

SIZE/COST FACTOR

Vessel size	GRT [1]	Average cost ($U.S.) per GRT
Extra Small	<3,001	0.25
Small	3,001–15,000	0.50
Medium	15,001–30,000	1.00
Large	30,001–60,000	1.50
Extra Large	>60,000	2.00

[1] Gross register tonnage in cubic feet, as shown in Lloyd's Register of Shipping.