Exhibit 2 shows the estimated annualized cost burden for the respondents. The Bureau of Labor Statistics reported that the average hourly wage for ''healthcare practitioner and technical occupations'' in the United States was $29.82 in May 2006. An estimate of $30 per hour allows for inflation and represents a conservative estimate of the wages of the respondents. Therefore, the total estimated cost burden for respondents is $12,030, based on the total estimated annualized burden of 401 hours.

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Form name	Number of respondents	Total burden hours	Average hourly wage rate *	Total cost burden
E-mail submission	41	21	$30	$630
Health care innovator interview—following e-mail submission	41	21	30	630
Health care innovator interview without e-mail submission	234	234	30	7,020
Annual follow-up interview	250	125	30	3,750
Total	566	401		12,030

* Based upon the average wages, ''National Compensation Survey: Occupational Wages in the United States, May 2006,'' U.S. Department of Labor, Bureau of Labor Statistics.

Estimated Annual Costs to the Federal Government

The total cost to the Government is approximately $3,349,560 over three years (on average, $1,116,520 per year). These costs cover the total editorial and content development processes associated with the project; which include developing an on-line authoring tool for preparing the profiles, identifying innovation leads, reviewing e-mail submissions, contacting the innovators, conducting interviews, preparing the draft profiles, securing innovator approval, and publishing the profiles on the Innovations Exchange Web site.

Request for Comments

In accordance with the above-cited Paperwork Reduction Act legislation, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ health care research, quality improvement and information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: August 12, 2008.

Carolyn M. Clancy,

Director.

[FR Doc. E8–19302 Filed 8–20–08; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: State Plan for the Temporary Assistance for Needy Families (TANF).

OMB No.: 0970–0145.

Description: The State plan is a mandatory statement submitted to the Secretary of the Department of Health and Human Services by the State. It consists of an outline of how the States TANF program will be administered and operated and certain required certifications by the States Chief Executive Officer. Its submittal triggers the States family assistance grant funding and it is used to provide the public with information about the program. If a State makes changes in its program, it must submit a State plan amendment.

Respondents: The 50 States, the District of Columbia, Guam, Puerto Rico and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Temporary Assistance to Needy Families State Plan Guidance	54	0.50	33	891

Estimated Total Annual Burden Hours: 891.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, *Fax:* 202–395–6974, *Attn:* Desk Officer for the Administration for Children and Families.

Dated: August 14, 2008.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. E8–19220 Filed 8–20–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0039]

Chloramine-T for Control of Bacterial Gill Disease in Freshwater-Reared Salmonids; Availability of Data

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of effectiveness and target animal safety data that may be used in support of a new animal drug application (NADA) or supplemental NADA for use of chloramine-T by immersion for the control of mortality in freshwater-reared salmonids due to bacterial gill disease. The data, contained in Public Master File (PMF) 5893, were compiled by the U.S. Department of the Interior, U.S. Fish and Wildlife Service, Aquatic Animal Drug Approval Partnership Program.

ADDRESSES: Submit NADAs or supplemental NADAs to the Document Control Unit (HFV–199), Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855.

FOR FURTHER INFORMATION CONTACT: Donald A. Prater, Center for Veterinary Medicine (HFV–130), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8343, e-mail: *donald.prater@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Chloramine-T used by immersion for control of mortality in freshwater-reared salmonids due to bacterial gill disease is a new animal drug under section 201(v) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 321(v)). As a new animal drug, chloramine-T is subject to section 512 of the act (21 U.S.C. 360b) which requires that its uses be the subject of an approved NADA or supplemental NADA. Fish are a minor species under § 514.1(d)(1)(ii) (21 CFR 514.1(d)(1)(ii)).

The U.S. Department of the Interior, U.S. Fish and Wildlife Service, Aquatic Animal Drug Approval Partnership Program, 4050 Bridger Canyon Rd., Bozeman, MT 59715, has provided effectiveness and target animal safety data for use of chloramine-T by immersion for control of mortality in freshwater-reared salmonids due to bacterial gill disease. These data are contained in PMF 5893.

Sponsors of NADAs or supplemental NADAs may, without further authorization, reference the PMF 5893 to support approval of an application filed under § 514.1(d). An NADA or supplemental NADA must include, in addition to reference to the PMF, animal drug labeling and other information needed for approval, such as: data concerning human food safety; and manufacturing methods, facilities, and controls. Persons desiring more information concerning PMF 5893 or requirements for approval of an NADA or supplemental NADA may contact the Center for Veterinary Medicine (see **FOR FURTHER INFORMATION CONTACT**).

In accordance with the freedom of information provisions of 21 CFR part 20, a summary of safety and effectiveness data provided in PMF 5893 to support approval of an application may be seen in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, from 9 a.m. to 4 p.m., Monday through Friday.

Dated: August 8, 2008.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. E8–19299 Filed 8–20–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Shared Instrumentation (S10) Review.

Date: September 8, 2008.

Time: 8:30 a.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency O'Hare, 9300 Bryn Mawr Avenue, Rosemont, IL 60018.

Contact Person: Barbara Whitmarsh, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2206, MSC 7890, Bethesda, MD 20892, (301) 435–4511, *whitmarshb@csr.nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Cell Biology Integrated Review Group Intercellular Interactions Study Section.

Date: September 22–23, 2008.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: George Washington University Inn, 824 New Hampshire Avenue, NW., Washington, DC 20037.

Contact Person: David Balasundaram, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5189, MSC 7840, Bethesda, MD 20892, 301–435–1022, *balasundaramd@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Applications Related to Dementia, Substance Abuse, or Behavioral Development.

Date: September 24, 2008.

Time: 2 p.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Ellen K. Schwartz, EDD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3168, MSC 7770, Bethesda, MD 20892, 301–435–0681, *schwarte@csr.nih.gov.*

Name of Committee: Bioengineering Sciences & Technologies Integrated Review Group; Biomaterials and Biointerfaces Study Section.

Date: September 25, 2008.

Time: 8 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: Doubletree Hotel Bethesda, 8120 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Ross D. Shonat, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5156, MSC 7849, Bethesda, MD 20892, 301–435–2786, *shonatr@csr.nih.gov.*

Name of Committee: Bioengineering Sciences & Technologies Integrated Review Group; Nanotechnology Study Section.

Date: September 30–October 1, 2008.

Time: 8:30 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Alexandria Old Town, 1767 King Street, Alexandria, VA 22314.

Contact Person: Joseph D. Mosca, PhD, Scientific Review Officer, Center for