TABLE 1—REGISTRATION FEES [1]

Attendee type	Advanced rate (March 13, 2013 to April 8, 2013)	Standard rate (April 9, 2013 to May 3, 2013)
Industry	$1,295	$1,495
Small Business (<100 employees)	900	1,000
Consultant	600	700
Startup Manufacturer	250	300
Academic	250	300
FDA/Government Employee	(2)	Free

[1] The following forms of payment will be accepted: American Express, Visa, Mastercard, and company checks.
[2] Free.

To register online for the public conference, please visit the ''Register Now'' link on the conference Web site at *http://www.XavierMedCon.com.* FDA has verified the Web site address, but is not responsible for subsequent changes to the Web site after this document publishes in the **Federal Register**.

To register by mail, please send your name, title, firm name, address, telephone and fax numbers, email, and payment information for the fee to Xavier University, Attention: Sue Bensman, 3800 Victory Pkwy., Cincinnati, OH 45207. An email will be sent confirming your registration.

Attendees are responsible for their own accommodations. The conference headquarter hotel is the Downtown Cincinnati Hilton Netherlands Plaza, 35 West Fifth St., Cincinnati, OH, 45202, 513–421–9100. Special Conference Block rates are available through April 9, 2013. To make reservations online, please visit the ''Venue & Logistics'' link at *http://www.XavierMedCon.com.*

If you need special accommodations due to a disability, please contact Marla Phillips (see Contact Persons) at least 7 days in advance of the conference.

SUPPLEMENTARY INFORMATION: The public conference helps fulfill the Department of Health and Human Services and FDA's important mission to protect the public health. The conference will provide those engaged in FDA-regulated medical devices (for humans) with information on the following topics:

- CDRH Future Vision and Strategy Keynote Address
- U.S. Congressman Erik Paulsen Keynote Dinner
- EU Regulations: New Regulations, Company Strategy, and Open Discussion Forum
- FDA Safety and Innovation Act
- Unique Device Identification
- Update from the Office of Device Evaluation
- Total Product Life Cycle: Interactive Workshop
- Pre-Submission Program and Meetings with the FDA
- 510(k): New FDA Guidance and Industry Regulations
- PMAs: New Guidance and Compliance Initiatives
- Software and Mobile Apps
- Combination Products
- Entering the EU Market and CE Mark Hot Topics
- Global Product Strategy
- Success in Central and South America
- FDA Inspectional Approach—Panel with Current FDA Investigators

FDA has made education of the drug and device manufacturing community a high priority to help ensure the quality of FDA-regulated drugs and devices. The conference helps to achieve objectives set forth in section 406 of the Food and Drug Administration Modernization Act of 1997 (21 U.S.C. 393), which includes working closely with stakeholders and maximizing the availability and clarity of information to stakeholders and the public. The conference also is consistent with the Small Business Regulatory Enforcement Fairness Act of 1996 (Pub. L. 104–121) by providing outreach activities by Government Agencies to small businesses.

Dated: March 8, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–05727 Filed 3–12–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-day Comment Request: Pediatric Palliative Care Campaign Pilot Survey

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Nursing Research (NINR), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on December 26, 2012, page 76053 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institute of Nursing Research (NINR), National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Ms. Adrienne Burroughs, Health Communications Specialist, Office of Communications and Public Liaison, NINR, NIH, Building 31, Room 5B10, 31 Center Drive, Bethesda, MD 20892 or call non-toll-free number (301) 496–0256 or Email your request, including your address to: *adrienne.burroughs@nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Pediatric Palliative Care Campaign Pilot Survey,

0925-New—National Institute of Nursing Research (NINR), National Institutes of Health (NIH).

Need and Use of Information Collection: NINR developed a Pediatric Palliative Care Campaign to address the communications challenges faced by health care providers who recommend and provide palliative care to pediatric populations. NINR is launching this effort to increase the use of palliative care for children living with serious illness or life-limiting conditions. The Pediatric Palliative Care Campaign Pilot Survey will assess the information and materials being disseminated as part of the Pediatric Palliative Care Campaign pilot. Survey findings will help (1) determine if the pilot campaign is effective, relevant, and useful to health care providers who recommend and provide palliative care to pediatric populations; (2) to better understand current perceptions, challenges, and information needs of health care providers when it comes to discussing pediatric palliative care so that information and materials can be refined; and (3) examine how effective the campaign pilot materials are in starting and continuing a pediatric palliative care conversation and addressing the communications needs of health care providers around this topic. This assessment will deliver strategic and actionable guidance for refining the campaign materials so that they can be used by a wider audience of health care providers.

OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 25.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Physicians	25	1	30/60	12.5
Nurses	25	1	30/60	12.5
Total	50	1	30/60	25

Dated: March 4, 2013.

Amanda Greene,

Science Evaluation Officer/Project Clearance Liaison, NINR, NIH.

[FR Doc. 2013–05774 Filed 3–12–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-day Comment Request: Early Career Reviewer Program Online Application System—Center for Scientific Review (CSR)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, Center for Scientific Review, the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) The quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

To Submit Comments and for Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Monica Basco, ECR Program, Center for Scientific Review, 6701 Rockledge Dr., Room 3220, Bethesda, MD 20892 or call non-toll-free number (301) 300–3839 or Email your request, including your address to: *CSRearlyCareerReviewer@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Collection: Early Career Reviewer Program Online Application System—Center for Scientific Review (CSR), 0925- New Information Collection Request), Center for Scientific Review (CSR), National Institutes of Health (NIH).

Need and Use of Information Collection: The Center for Scientific Review (CSR) is the portal for NIH grant applications and their review for scientific merit. Our mission is to see that NIH grant applications receive fair, independent, expert, and timely reviews—free from inappropriate influences—so NIH can fund the most promising research. To accomplish this goal, Scientific Review Officers (SRO) form study sections consisting of scientists who have the technical and scientific expertise to evaluate the merit of grant applications. The CSR Early Career Reviewer (ECR) program was developed to identify and train qualified scientists who are early in their scientific careers and who have not had prior CSR review experience. Currently, the application process involves repeated email interactions with potential applicants and manual management of information. To make the application process more efficient for applicants and for CSR staff, we are working with the Information Management Branch at CSR to develop online application software which includes the collection of applicants' names, contact information, and professional CV. This PRA clearance request is to develop online application software for ECR program applicants.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 650.