TABLE 1—LIST OF SAFETY AND EFFECTIVENESS SUMMARIES FOR APPROVED PMAS MADE AVAILABLE FROM OCTOBER 1, 2008, THROUGH DECEMBER 31, 2008.

PMA No./Docket No.	Applicant	TRADE NAME	Approval Date
P070015 FDA–2008–M–0535	Abbott Vascular Inc.	XIENCE V EVEROLIMUS ELUTING CORONARY STENT SYSTEM & PROMUS ELUTING CORONARY STENT SYSTEM	July 2, 2008
P030025 (S28) FDA–2008–M–0547	Boston Scientific Corp.	TAXUS EXPRESS2 PACLITAXEL ELUTING CORONARY STENT SYSTEM	September 24, 2008
P080004 FDA–2008–M–0536	Hoya Surgical Optics, Inc.	HOYA ISPHERIC MODEL YA–60BB INTRAOCULAR LENS	September 26, 2008
H070004 FDA–2008–M–0563	Levitronix, LLC	LEVITRONIX CENTRIMAG RIGHT VENTRICULAR ASSIST SYSTEM (RVAS)	October 7, 2008
P060008 FDA–2008–M–0593	Boston Scientific Corp.	TAXUS LIBERTE' PACLITAXEL ELUTING CORONARY STENT SYSTEM	October 10, 2008
P050029 FDA–2008–M–0601	Stereotaxis, Inc.	HELIOS II ABLATION CATHETER	October 10, 2008
H040004 FDA–2008–M–0562	Medtronic Sofamor Danek USA, Inc.	INFUSE/MASTERGRAFT POSTEROLATERAL REVISION DEVICE	October 10, 2008
P050019 FDA–2008–M–0596	Boston Scientific Corp.	CAROTID WALLSTENT MONORAIL ENDOPROSTHESIS	October 23, 2008
H060002 FDA–2008–M–0579	Spiration, Inc.	IBV VALVE SYSTEM	October 24, 2008
P060025 FDA–2008–M–0594	ATS Medical, Inc.	ATS 3F AORTIC BIOPROSTHESIS	October 30, 2008
P080011 FDA–2008–M–0608	Coopervision Manufacturing, Ltd.	BIOFINITY COMFILCON A (EXTENDED WEAR SOFT CONTACT LENSES)	November 19, 2008
P080007 FDA–2008–M–0645	Bard Peripheral Vascular Inc.	BARD E-LUMINEXX VASCULAR STENT	December 4, 2008
P060006 FDA–2008–M–0646	Boston Scientific Corp.	BOSTON SCIENTIFIC EXPRESS SD RENAL MONORAIL PREMOUNTED STENT SYSTEM	December 11, 2008

II. Electronic Access

Persons with access to the Internet may obtain the documents at *http://www.fda.gov/cdrh/pmapage.html*.

Dated: March 10, 2009.

Daniel G. Schultz,

Director, Center for Devices and Radiological Health.

[FR Doc. E9–6026 Filed 3–18–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Simulations for Drug Related Science Education

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute on Drug Abuse (NIDA), 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 June 26, 2008, (Vol. 73 No. 124, page 36337) 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 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 November 15, 2008, unless it displays a currently valid OMB control number.

Proposed Collection: Title: Simulations for Drug Related Science Education. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* This is a request for a one-time clearance to evaluate an interactive multimedia module developed by *ArchieMD*. This evaluation seeks to determine whether the multimedia module *Archie MD: The Science of Drugs* (1) Increases students' knowledge in brain and heart biology and the effects drugs have on the body (2) Increases positive attitudes towards science education for high school students (3) Reinforce or instill negative attitudes towards substance abuse. In order to test the effectiveness of the interactive multimedia module, data will be collected in the form of pre and post test surveys from 10th and 11th grade high school students utilizing the developed module. The findings will provide valuable information regarding information pertaining to the use of interactive multimedia educational modules in high school science classrooms and their ability to increase knowledge and change attitudes and perceptions.

Frequency of Response: 4. *Affected Public:* High school students engaged with the *ArchieMD: The Science of*

Drugs program. *Type of Respondent:* Participants will include high school students enrolled in the tenth and eleventh grade. *Estimated Total Annual Number of Respondents:* 360. *Estimated Number of Responses per Respondent:* 4. *Average Burden Hours per Response:* One high school period lasting 50 minutes. *Estimated Total Annual Burden Hours Requested:* 1199.95. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report. The estimated annualized burden is summarized below.

Type of respondents	Number of respondents	Frequency of response	Average burden hours per response	Estimated total burden hours requested
Participants—High School Students	360	4	.8333	1199.95
Total	360	4	.8333	1199.95

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (2) the accuracy of the agency's estimate of the burden of the proposed collection of information; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

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: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. Cathrine Sasek, Coordinator, Science Education Program, Office of Science Policy and Communications, National Institute on Drug Abuse, 6001 Executive Blvd, Room 5237, Bethesda, MD 20892, or call non-toll-free number (301) 443–6071; fax (301) 443–6277; or by e-mail to *csasek@nida.nih.gov*.

Comments 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.

Dated: March 12, 2009.

Mary Affeldt,

Associate Director for Management, National Institute on Drug Abuse.

[FR Doc. E9–6037 Filed 3–18–09; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Survey of NHLBI Constituents' Health Information Needs and Preferred Formats

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, the National Heart, Lung and Blood Institute (NHLBI), 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.

Proposed Collection: Title: Survey of NHLBI Constituents' Health Information Needs and Preferred Formats. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* The purpose of this survey is to obtain information from NHLBI constituents (health professionals, patients and their families, and the general public) for the purpose of evaluating their health information and education needs and format preferences. The Consumer Services Team (CST) will use the data collected in this survey to create a 3-year Strategic Plan. The findings from the survey, described in the Strategic Plan, will be used to develop new health information materials for NHLBI constituents and to revise materials currently in the Institute's portfolio. *Frequency of Response:* Once every 3 years. *Affected Public:* Individuals. *Type of Respondents:* Individuals who have been consumers of NHLBI information within the past 3 years. The annual reporting burden is as follows: *Estimated Number of Respondents:* 2,450; *Estimated Number of Responses per Respondent:* 1; *Average Burden Hours Per Response:* 0.2; and *Estimated Total Annual Burden Hours Requested:* 162. The annualized cost to respondents is estimated at: $3,518.62. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
General Public	1,075	0.33	0.2	71
Private Companies	332	0.33	0.2	22
Public Sector Groups	332	0.33	0.2	22
Health Professionals	711	0.33	0.2	47
Totals	2,450	.....	.....	162

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on 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