by prospectively studying children who were born with this potentially disabling condition. We estimate to enroll approximately 40 parents with a child with Spina Bifida ages 3-, 4-, or 5-years of age, and 20 of the children of these forty parents. The data to be collected will relate to medical concerns prevalent among individuals with Spina Bifida in the areas of neurology/neurosurgery, urology, and orthopedics; development and learning; nutrition and physical growth; mobility and functioning; general health; and family demographics. Families interested in participating can choose between participating in a phone survey (no more than 40 minutes) or an in-person assessment (no more than 2 hrs). For families who participate in the in-person assessment, (estimated to be twenty of the forty families); the child will also be invited to participate in a child-appropriate assessment. Data will also be collected on the actual recruitment process. Results from the project will be evaluated and disseminated to provide guidance for states that are interested in following children with Spina Bifida prospectively. The proposed project is the initial step to document the development, the health status, and the onset of complications among children with SB in order that effective interventions may be identified that will ameliorate the course of this complex, multi-system condition. Long-term results will help determine if it would be beneficial to systematically screen children with Spina Bifida for certain health related, educational and developmental problems that these children are at an increased risk of experiencing and at what age such a screening should be performed.

There will be no cost to the respondents other than their time.

ESTIMATE OF ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in minutes)	Total burden hours
Parents (phone survey)	20	1	40/60	13
Parents (in-person assessment)	20	1	2	40
Child (in-person assessment)	20	1	1	20
Total				73

Dated: January 25, 2008.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E8–1993 Filed 2–4–08; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 72 FR 67308, dated November 28, 2007) is amended to reflect the title change for the Division of Nutrition, Physical Activity, and Obesity Prevention, National Center for Chronic Disease Prevention and Health Promotion, Coordinating Center for Health Promotion, Centers for Disease Control and Prevention.

Section C–B, Organization and Functions, is hereby amended as follows:

Delete in its entirety the title for the *Division of Nutrition, Physical Activity, and Obesity Prevention (CUCH)* and insert the Division of Nutrition, Physical Activity, and Obesity (CUCH).

Dated: January 28, 2008.

Joseph Henderson, M.P.A.,

Acting Chief Operating Officer, Centers for Disease Control and Prevention (CDC).

[FR Doc. 08–486 Filed 2–4–08; 8:45 am]

BILLING CODE 4160–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0051] (formerly Docket No. 2007N–0422)

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Application for Participation in the Medical Device Fellowship Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by March 6, 2008.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974, or e-mailed to *baguilar@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0551. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley Jr.,Office of the Chief Information Officer (HFA–250), Food and Drug Administration,5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Application for Participation in the Medical Device Fellowship Program; (OMB Control Number 0910–0551)—Extension

Sections 1104, 1302, 3301, 3304, 3320, 3361, 3393, and 3394 of title 5 of the United States Code, authorize Federal agencies to rate applicants for Federal jobs. Collecting applications for the Medical Device Fellowship Program will allow FDA's Center for Devices and Radiological Health (CDRH) to easily and efficiently elicit and review

information from students and health care professionals who are interested in becoming involved in CDRH activities. The process will reduce the time and cost of submitting written documentation to the agency and lessen the likelihood of applications being misrouted within the agency mail system. It will assist the agency in promoting and protecting the public health by encouraging outside persons to share their expertise with CDRH.

In the **Federal Register** of November 9, 2007 (72 FR 63614), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

FDA estimate of the burden for this collection of information is as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

5 U.S.C. Section/ FDA Form No.	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
1104, 1302, 3301, 3304, 3320, 3361, 3393, 3394/ Form No. 3608	250	1	250	1	250

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA based these estimates on the number of inquiries that have been received concerning the program and the number of requests for application forms over the past 3 years.

Dated: January 30, 2008.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E8–2068 Filed 2–4–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0048] (formerly Docket No. 2007N–0182)

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Information Program on Clinical Trials for Serious and Life-Threatening Diseases: Maintaining a Databank

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Information Program on Clinical Trials for Serious and Life-Threatening Diseases: Maintaining a Databank'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of October 19, 2007 (72 FR 59295), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0459. The approval expires on January 31, 2011. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: January 30, 2008.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E8–2076 Filed 2–4–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0050]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices; Device Tracking

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements for the tracking of medical devices.

DATES: Submit written or electronic comments on the collection of information by April 7, 2008.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration,5600 Fishers Lane, Rockville, MD 20857, 301 827–1472.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites