Dated: August 12, 2002.

Sandra R. Manning,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 02–20947 Filed 8–16–02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Proposed Projects

Title: National Survey of Child and Adolescent Well-Being.

OMB No.: 0970–0202.

Description: This longitudinal survey provides national estimates on the characteristics related to children and fami8lies who enter the child welfare system. It has collected data from a cohort of 6,100 children who entered the child welfare system as a result of a CPS investigation between October 1999 and April 2001. Data were collected from the children themselves, their caregivers, their teachers, and their caseworkers at baseline, with follow-ups at 12 and 18 months post-baseline. The current request is to pursue a 36-month follow-up, essentially replicating the measure that were used at baseline and at the 18-month follow-up.

Respondents: Children who are clients of the child welfare system, their primary caregivers, caseworkers, and teachers.

Annual Burden Estimates:

Instrument	Responses	Number of responses per respondent	Average burden hours per respondent	Total burden hours
Child interview	5,491	1	1.63	8,950
Caregiver interview	5,491	1	1.50	8,237
Caseworker Interview	2,366	1	0.80	1,893
Caseworker Interview	2,491	1	0.75	1,868

Estimated Total Annual Burden Hours: 20,948.

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.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 650 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 Act, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Dated: August 13, 2002.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 02–20936 Filed 8–16–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0355]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Device Recall Authority

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 information collection, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements for medical device recall authority.

DATES: Submit written or electronic comments on the collection of information by October 18, 2002.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (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: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

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 comments on: (1) Whether the proposed collection of information is necessary